Title: Clinical decision making and mental health service use in people with severe mental illness across Europe

Disclosures and acknowledgments.

The authors have no conflict of interest to declare.

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Abstract

Objective: This study aims to explore relationships between preferred and experienced clinical decision making with service use, and associated costs, by people with severe mental illness.

Methods: Prospective observational study of mental healthcare in six European countries: Germany, UK, Italy Hungary, Denmark and Switzerland. Patients (N = 588) and treating clinicians (N = 213) reported preferred and experienced decision making at baseline using the Clinical Decision Making Style Scale (CDMS) and the Clinical Decision Involvement and Satisfaction Scale (CDIS). Retrospective service use was assessed with the Client Socio-Demographic and Service Receipt Inventory (CSSRI-EU) at baseline and 12-month follow-up. Negative binominal regression analyses examined the effects of CDMS and CDIS on service use and inpatient costs at baseline and multilevel models examined these relationships over time.

Results: At baseline, staff and patient preferences for active decision making and low patient satisfaction with experienced decisions were associated with longer hospital admissions and higher costs. Patient preferences for active decision making predicted increases in hospital admissions (b = .236, p = .043) over 12 months and cost increases were predicted by low patient satisfaction (b = 4803, p = .005). Decision making was unrelated to medication, outpatient, or community service use.

Conclusions: Decision making is related to inpatient service use and associated costs by people with severe mental illness. A preference for shared decision making may reduce healthcare costs via a reduction in inpatient admissions. Patient satisfaction with decisions is a crucial predictor of healthcare costs; therefore, clinicians should maximize patient satisfaction with decision making.

Key words: clinical decision making, severe mental illness, mental health care, service use, costs
Introduction

There is widespread agreement that shared, rather than passive, clinical decision making between patients and staff is an ethical obligation (1–3). A collaborative process of decision making contributes to high-quality, patient-centered healthcare (4), with some evidence indicating that shared decision making might have positive effects on treatment outcomes (1,5,6). There is also growing evidence suggesting that, in medical settings, shared decision making may be associated with cost-effectiveness (7), by reducing hospital admissions and other associated medical costs (8).

Although the extant body of literature exploring decision making has largely focused on physical health conditions, decision making is a growing area of interest in psychiatric settings, especially in the treatment of severe mental illness (9–11) with psychiatric patients shown to be capable of (3) and expressing a preference for shared decision making (12–14,1,15). However it remains largely unknown if a preference for or experiences of shared decision making impacts upon service use. It has been suggested that higher involvement of mental health patients in decision making might increase their engagement with services (16), although empirical evidence remains equivocal (1). Likewise, some evidence indicates that satisfaction with decision making has a positive relationship with medication adherence (17), yet other studies have found no such relationship (18). Although there is limited evidence that shared decision making may reduce physical healthcare costs through the reduction of utilizing ineffective or undesirable treatments (19), whether such cost-effectiveness is seen in mental health care remains unknown.

Despite the potential links between decision making and service use and satisfaction with services, research exploring decision making and its impact on adherence, engagement and service use in mental health remains scant (20), especially in real world psychiatric settings (6). The extent to which experiences of CDM and preferred CDM impact service use and the associated costs remains largely unknown. Therefore, the current study aims to examine decision making in routine psychiatric care for people with severe mental illness through an exploration of the relationship between preferred and experienced CDM with service use. Specifically, this study examines the following questions: a) Do CDM preferences regarding participation in decision making influence service use and costs? b) Does preference for the level of information impact service use? c) Does experienced level of
involvement in decision making impact service use? d) Does satisfaction with experienced clinical decisions impact service use?

**Method**

**Design**

The study was undertaken as part of the European multicenter study, “Clinical decision making and outcome in routine care for people with severe mental illness” (CEDAR, ISRCTN75841675) (21); a prospective observational study of routine psychiatric care. Patients and staff were recruited from outpatient/community mental health services in six European countries (Germany, UK, Italy, Hungary, Denmark, and Switzerland) between November 2009 and December 2010. Ethical approval was obtained from all participating centers. We used CDM data collected at baseline and service use data that were collected at baseline and 12-month follow up.

**Participants**

Participants who presented with any psychiatric diagnosis - established by case notes or staff communication, based on SCID criteria (22,23) - were recruited from six European centers. Inclusion criteria were (a) aged 18-60, (b) presence of severe mental illness (Threshold Assessment Grid (24) ≥ 5 points and illness duration ≥ 2 year), (c) expected contact with mental health services for one year, (d) sufficient language proficiency, and (e) capacity to provide informed consent. Exclusion criteria were (a) main diagnosis of learning disability, dementia, substance misuse or organic brain disorder, (b) severe cognitive impairment, or (c) treatment by forensic mental health services. At baseline, patients identified a key professional involved in their care, who was then approached for study participation. Patients and professionals who provided informed consent following an explanation of the study protocol were included in the study (see Puschner et al. (21) for further information on study procedures).
Measures

**Client Sociodemographic and Service Use Inventory (CSSRI-EU).** The CSSRI-EU (25) provides self-report information in five areas: (a) sociodemographic data including ethnicity and education level, (b) living situation, (c) employment and income, (d) service receipt such as inpatient and outpatient hospital services, primary and secondary community care contacts, and (e) medication use. For pragmatic reasons, the data on medication was assessed using a shortened version of the CSSRI (Paul McCrone, personal communication), with patients being asked to indicate type (psychotropic or non-psychotropic) and number of medications (while the original version also asks for brand name and dose). Community and outpatient services for the prior three months, and number of medications used in the previous one month, were collected. Number and duration of inpatient stays was collected for the previous 12 months. Total costs of inpatient days were calculated by allocating unit costs based on average costs per day of inpatient stays in each participating center during the observation period.

**Clinical Decision Making Style Scale (CDMS).** The CDMS (27) measures preferences for CDM. Identically structured parallel versions of the CDMS have been designed for staff and patients. The CDMS is divided into three sections and yields two subscales: Participation in Decision Making (PD) and Information (IN). PD is measured by two sections: a) agreement on a 5-point Likert scale with six items such as “important decisions should be made by the clinician in charge and not by me” and b) ratings on nine items regarding if the decision should be made by the clinician, patient or shared, based on three vignettes in the areas of work, medication and side effects. The PD subscale score is then classified into one of three categories: a preference for active, shared, or passive decision making. The IN subscale is measured by agreement on a 5-point Likert scale with six items regarding the need of the patient to receive information. Scores are categorized into preference for either high, moderate, or low levels of information provision.
Clinical Decision Making Involvement and Satisfaction Scale (CDIS) (26). This instrument measures involvement and satisfaction with a recently experienced clinical decision, as rated by both patient and staff. The Involvement (CDIS-INV) subscale is assessed through a single item rating the extent to which the decision was shared, passive or actively made by the patient (rated on a 5-point scale). The Satisfaction (CDIS-SAT) subscale is assessed by level of agreement on a 5-point Likert scale with six items regarding a) being informed, b) making the best decision, c) consistency with personal values, d) expectation of implementing the decision, e) whether this was the best decision to make, and f) overall satisfaction. The satisfaction score is then classified into three categories: high, moderate and low. CEDAR measures and scoring information can be downloaded at www.cedar-net.eu/instruments.

Analysis

Data were analyzed using SPSS for Windows 23.0. Comparisons of service use, as assessed by the CSSRI-EU, at baseline and follow-up were conducted using non-parametric paired Wilcoxon rank tests, with Bonferroni adjustment for multiple analyses. Costs of inpatient admissions at baseline and follow-up were assessed via paired t-tests. Predictors of baseline service use were assessed via negative binomial regression analyses, entering all baseline staff-rated and patient-rated subscales of the CDMS and CDIS as predictors, after controlling for age, gender, marital status, study site, diagnosis and duration of illness. To assess changes in service use over time, multilevel models were used with a random slope and intercept; controlling for the same covariates.

Results

A total of 588 psychiatric outpatients from the six centers participated. Mean age of participants was 41.7 years (SD = 10.7). The majority of participants were unmarried (n = 439, 75%), unemployed (n = 478, 81%) and female (n = 307, 52%). The most common DSM-IV diagnoses were psychotic disorders (n = 269, 46%) and mood disorders (n = 200, 34%), and
the average duration of illness was 12.5 years (SD = 9.3). Each participant had an allocated staff member. Paired data for the 588 patients were obtained from staff members (N = 213); with some staff members providing data for multiple patients. Staff members had a mean age of 45.9 years (SD = 10.6) and 62% were female (n = 127). Of those participating, 37% were psychiatrists (n = 75), 9% psychologists (n = 19), 5% social workers (n = 11), whilst 49% were from another profession within mental health services (n = 100). On average, staff had worked in mental health for 15.0 years (SD = 9.7).

At baseline, the majority of both patients and staff expressed a preference on the CDMS-PD for shared (71%, 54% respectively) rather than active (8%, 16%) or passive (22%, 30%) decision making. The majority of patients reported a preference on the CDMS-IN for receiving high levels of information regarding decisions, whereas the majority of staff preferred patients to receive a moderate amount (high 61%, 35%; mod 35%, 57%; low 4%, 8%). When asked at baseline about the most recently experienced clinical decision on the CDIS-INV, nearly half of all patients and staff reported that the decision had been shared (50%, 47%; passive 27%, 23%; active 24%, 30%). The majority of staff and patients reported being moderately or highly satisfied with the decision on the CDIS-SAT (high 52%, 45%; mod 43%, 51%; low 6%, 4%).

Community contacts and day services were the most commonly used services at both baseline and follow-up, with a total of 201 inpatient admissions across the sample at baseline and 137 by follow-up (see Table 1). Comparison of self-reported service use showed that number and length of inpatient stays significantly decreased between baseline and follow-up. Likewise, total costs of inpatient admissions significantly decreased from €2 512 330 at baseline to €807 803 at follow up \(t = 3.82, \text{ df } =519, \ p >.001\). No significant changes were observed for the number of outpatient, day service and community contacts, nor for number of medications prescribed.

###Table 1 about here###

**Relationship between decision-making and baseline service use**

At baseline, a patient-rated preference on the CDMS-PD for an active style was associated with a higher number of inpatient stays than preferences for passive of shared style (see Table 2).

###Table 2 here ###
The duration and costs of such inpatient admissions were significantly predicted by all subscales, both staff- and patient-rated. Patient-rated CDMS preferences for passive or shared CDM style were associated with shorter admissions than preferences for active CDM style and, relatedly, with lower costs compared with preferences for active CDM. On the staff-rated CDMS-PD, a preference for shared CDM style predicted slightly longer admissions, and preferences for a passive style were related to decreased costs. A preference for low levels of information provision on the CDMS-IN (both staff- and patient-rated) predicted shorter admissions and lower costs than high CDMS-IN. Similarly, both staff- and patient-reported passive involvement in a recent decision on the CDIS-INV predicted fewer days of inpatient admissions than active involvement, and passive and shared CDIS-INV predicted lower costs. Patient-rated low satisfaction on the CDIS-SAT predicted an increased number of admission days and increased costs, whereas a higher number of days and costs were associated with moderate staff-rated satisfaction. The number of outpatient contacts, day services, community contacts, and prescribed medications were not associated with the CDMS or CDIS subscales.

Relationship between decision-making and service use over time

Significant changes in service use were further examined to identify predictors thereof. Number of inpatient admissions was significantly predicted by both patient- and staff-rated CDMS-PD. An increase in the number of stays over time was predicted by patient preference for active decision making ($b = .236$, $t = 2.03$, df = 353, $p = .043$) and a staff preference for shared decision making ($b = .181$, $t = 2.21$, df = 353, $p = .028$). A longer length of inpatient stay was predicted by staff-rated preference for shared decision making compared with a patient preference for a passive style ($b = 5.49$, $t = 2.17$, df = 706, $p = .030$). The change in costs over one year was predicted by the patient-rated CDIS-SAT, with low satisfaction associated with an increase in costs ($b = 4803$, $t = 2.83$, df = 752, $p = .005$).

Discussion
This study contributes to the limited exploration of clinical decision making and its association with mental health service use amongst people with SMI. Decision making preferences and experiences were associated with number and duration of inpatient admissions and associated costs, both at baseline and with changes over 12-months. Measures of decision making were, however, unrelated to use of medications, and community, day and outpatient services.

Consistent with a growing body of evidence suggesting that, in medical settings, decision making style may be associated with cost-effectiveness (7,8), the present findings also suggest that aspects of decision making are related to service use and costs. Specifically, both preferences for and experiences of passive and shared, as opposed to active, decision making were associated with reduced hospital admissions and costs. Although reduced costs do not equate to cost-effectiveness, such findings are in line with the observations from physical health settings (8) that shared decision making might improve cost-effectiveness, possibly via improved treatment and thus a reduction in the need for hospital admissions (4,7,8). This finding may also reflect shared factors that influence both decision making and treatment outcomes. Those who are less unwell are more likely to express a preference for shared decision making (1, 14) and, concomitantly, are less likely to have inpatient admissions, especially lengthy ones. Additionally, it has previously been reported that patients’ belief of their own decisional capacity influences CDM preferences (31). Thus, preferences for participation in decision making and level of involvement in decisions may be a marker for other factors such as insight, understanding about the illness, and illness severity, which are related to inpatient stays and costs. Our results showed that those preferring and experiencing active CDM had more admissions, thus active, as opposed to passive or shared CDM may be a marker for those who are less well. The extent to which the relationship between decision making and admission costs is mediated by insight and clinical presentation would be a beneficial avenue for future research. More rigorously designed studies are needed to ascertain if decision making has a causal association with reducing admissions and costs.

A low patient satisfaction with experienced decision making predicted longer admissions and higher costs at baseline, as well as an increase in costs over time; highlighting the importance of patient satisfaction with their experienced decisions. This finding is in line with earlier research suggesting that patient satisfaction with explanations offered by service providers is associated with service use in people with severe mental illness (29). Whether satisfaction with actual decision making
independently drives the association with costs and admissions, or whether dissatisfaction is associated with poorer wellbeing and greater illness severity and therefore higher costs remains unclear. Although evidence that decision making interventions improve satisfaction is currently limited, this finding provides further support that the ongoing development of strategies to maximize patient satisfaction with CDM experiences (e.g., 33,34) may be beneficial in reducing admissions. Relatedly, the extent to which differential preferences of patient and treating clinician impact upon experienced CDM, especially patient satisfaction, would be a value avenue for future exploration. Given the relationship between satisfaction and service use, staff awareness of patient preferences and expectations would likely be valuable to assist in enhancing patient satisfaction.

The study lends support for the assertion that there is no relationship between preferred and experienced CDM and medication use in outpatient mental health settings (18), contrasting previous findings showing satisfaction is related to medication adherence (17). Additionally, despite previous evidence suggesting that involvement in decision making might increase engagement with community care services (16), in the present study, the use of outpatient and community services was unrelated to CDM preferences and experiences. Although engagement was not directly studied, outpatient service use may serve as a proxy indicator of engagement. Thus our results suggest that decision making, at least in those with SMI, may be largely unrelated to engagement with outpatient services. Such a finding may reflect that other factors contribute more to engagement with services, or may also reflect that those participating in the study and completing bimonthly questionnaires are already more engaged.

Limitations

Some limitations are noted in the present study. Firstly, the majority of the sample reported being highly satisfied with their experienced CDM and this might have limited the range of findings regarding relationships between satisfaction and outcomes. Secondly, the experienced involvement in decision making was assessed using a single item, possibly limiting the conclusions that can be drawn with service use. Relatedly, the scales used were developed and validated within the study sample. As such, the ability of the scales to differentiate between CDM experiences and preferences across settings and patient groups
cannot be certain. Additionally, only costs associated with inpatient admissions were assessed. Multivariate analyses controlled for a range of demographic and illness-related covariates, however, models are not exhaustive and additional factors which could impact the relationship between decision making and service use might have been overlooked. The sample examined was heterogeneous, with mixed diagnoses and co-morbidities and patients were treated in different clinics by different clinicians. The staff also came from a diversity of professional backgrounds. Whilst heterogeneity might limit the conclusions that can be drawn about the relationship between CDM preferences and experiences and service use, the real world nature of the study increases the generalizability of the findings and builds on the recommendation for naturalistic CDM research (2,4).

**Conclusions**

Findings highlight that several aspects of decision making are associated with inpatient service use, but not with use of other services. The extent to which decision making preferences and experiences are a marker for wellbeing and insight, or independently drive the associations with service use requires ongoing exploration. Clinicians should ensure patients are satisfied with their experiences of decision making. Decision making interventions might be fruitful for enhancing treatment provision and, ultimately cost-effectiveness, in mental health services.
References


Table 1. Self-reported service use at baseline and 12-month follow up (N = 578)

<table>
<thead>
<tr>
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<th>Baseline</th>
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<th>12-month follow up</th>
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<tbody>
<tr>
<td></td>
<td>Total</td>
<td>Md</td>
<td>IQR</td>
<td>Total</td>
</tr>
<tr>
<td>Inpatient days (previous year)</td>
<td>6547</td>
<td>0.0</td>
<td>0.0-1.0</td>
<td>2871</td>
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<td>Inpatient stays (previous year)</td>
<td>201</td>
<td>0.0</td>
<td>0.0-1.0</td>
<td>137</td>
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<td>Outpatient visits (previous 3 months)</td>
<td>2716</td>
<td>1.0</td>
<td>1.0-6.0</td>
<td>3165</td>
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<td>Day services (previous 3 months)</td>
<td>2683</td>
<td>0.0</td>
<td>0.0-1.0</td>
<td>2236</td>
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<td>Community contacts (previous 3 months)</td>
<td>5769</td>
<td>5.0</td>
<td>1.0-14.0</td>
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<td>Psychotropic medications (previous month)</td>
<td>1555</td>
<td>2.0</td>
<td>2.0-4.0</td>
<td>1332</td>
</tr>
</tbody>
</table>

Notes: Md = median; IQR = interquartile range; <sup>a</sup>Wilcoxon signed-rank test; <sup>b</sup>Bonferroni adjusted p = 0.05/6 = 0.008
### Table 2: Predictors of inpatient stays, days and costs at baseline

<table>
<thead>
<tr>
<th></th>
<th>Inpatient stays&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Inpatient days&lt;sup&gt;b&lt;/sup&gt;</th>
<th>Costs of inpatient days&lt;sup&gt;c&lt;/sup&gt;</th>
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<td></td>
<td>b</td>
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<td>OR</td>
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<tr>
<td><strong>Patient-rated</strong></td>
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<tr>
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<td>.233</td>
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<tr>
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<td>.884</td>
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<tr>
<td>CDIS SAT low&lt;sup&gt;§&lt;/sup&gt;</td>
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<td>.453</td>
<td>.190</td>
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<td>CDIS SAT moderate&lt;sup&gt;§&lt;/sup&gt;</td>
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**Notes:** All models controlled for age, gender, marital status, illness duration, DSM IV disorder, and study site; a: $\chi^2(23) = 57.6$, p<.001; b: $\chi^2(23) = 414.5$, p<.001; c: $\chi^2(22) = 1012$, p<.001; ^ reference group= active CDM; ◊ reference group = high information; * reference group= high involvement; § reference group= high satisfaction.